



Scan to download a copy of this presentation

# IMPACT OF ERENUMAB ON ACUTE MEDICATION USAGE AND HEALTHCARE RESOURCE UTILIZATION AMONG MIGRAINE PATIENTS: A US CLAIMS DATABASE STUDY

**Stewart J Tepper<sup>1</sup>, Juanzhi Fang<sup>2</sup>, Pamela Vo<sup>3</sup>, Ahmad Abdrabboh<sup>2</sup>, Matias Ferraris<sup>3</sup>, Ying Shen<sup>4</sup>, Lujia Zhou<sup>4</sup>, Mrudula Glassberg<sup>2</sup>**

<sup>1</sup>Geisel School of Medicine at Dartmouth, Hanover, NH, USA; <sup>2</sup>Novartis Pharmaceuticals Corporation, East Hanover, NJ, USA; <sup>3</sup>Novartis Pharma AG, Basel, CH-4002, Switzerland; <sup>4</sup>KMK Consulting Inc., Morristown, NJ, USA

**Disclosures:** **Stewart Tepper** – Salary: Dartmouth-Hitchcock Medical Center, American Headache Society; Stock options: Nocira, Percept; CME honoraria: American Academy of Neurology, American Headache Society, Cleveland Clinic Foundation, Diamond Headache Clinic, Elsevier, Forefront Collaborative, Hamilton General Hospital, Ontario, Canada, Headache Cooperative of New England, Henry Ford Hospital, Detroit, Inova, Medical Learning Institute Peerview, Miller Medical Communications, North American Center for CME, Physicians' Education Resource, Rockpointe, WebMD/Medscape; **Juanzhi Fang, Pamela Vo, Ahmad Abdrabboh, Mrudula Glassberg and Matias Ferraris** – employees of, and own stocks in, Novartis; **Ying Shen and Lujia Zhou** – employees of KMK Consulting Inc.

Migraine Trust Virtual Symposium, 3–9 October, 2020

This study is supported by Novartis Pharma AG, Basel, Switzerland. Erenumab is co-developed by Amgen and Novartis.

Medical writing support was provided by Bronwyn Boyes of Novartis Ireland, Ltd.

The final responsibility for the content lies with the authors.



## Background and Methods

- This was a retrospective non-interventional observation analysis using Optum's de-identified Clinformatics® Data Mart (CDM) database
- Adult migraine patients initiating erenumab between 1 May 2018 and 30 September 2019 were identified

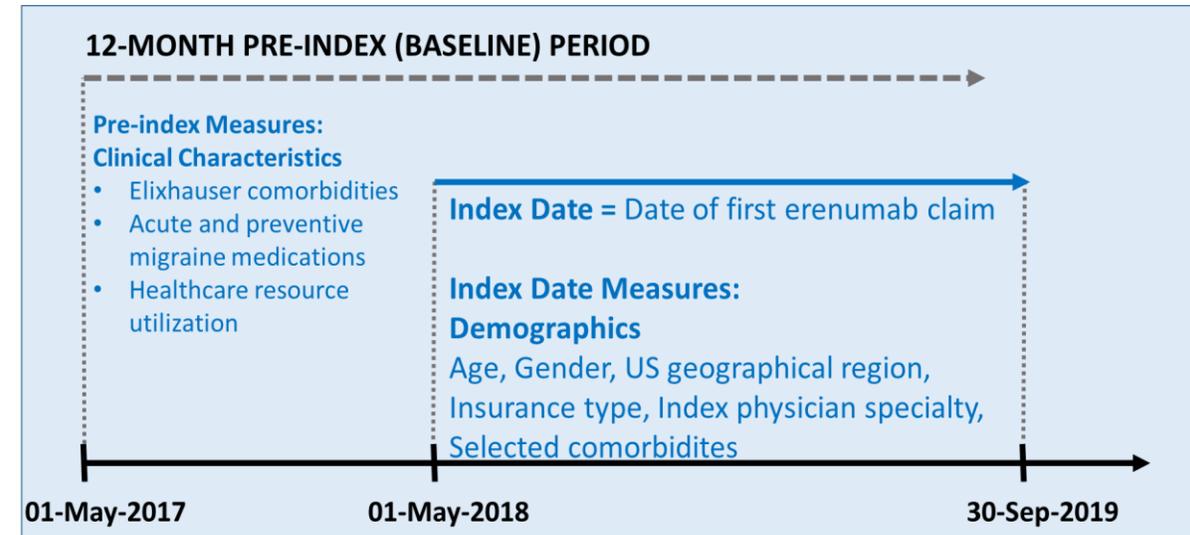
### Outcomes

- 6-month acute medication usage before and after erenumab treatment\*
- 6-month HCRU (including neurologist/headache specialist visits) before and after erenumab treatment
- Composite endpoint of outpatient visit with a diagnosis of migraine and an associated acute medication claim, hospital admission or emergency room visit with a primary diagnosis for migraine

\*Note that use of non-migraine specific acute medications (NSAIDs, opioids, and barbiturates) required a migraine diagnosis on or before seven days of the medication claim

CDM, Clinformatics® Data Mart; CI, confidence interval; HCRU, health care resource utilization; US, United States of America.

Figure 1: Study design



### Statistical analysis

- Descriptive analysis was performed
- Negative binomial model with repeated measure for count variables, such as the number of visits and claims, with rate ratios and 95% confidence interval (CI) calculated
- McNemar test for dichotomous variables

# Results

**Table 1. Demographics and 12-month pre-index characteristics of the population analyzed**

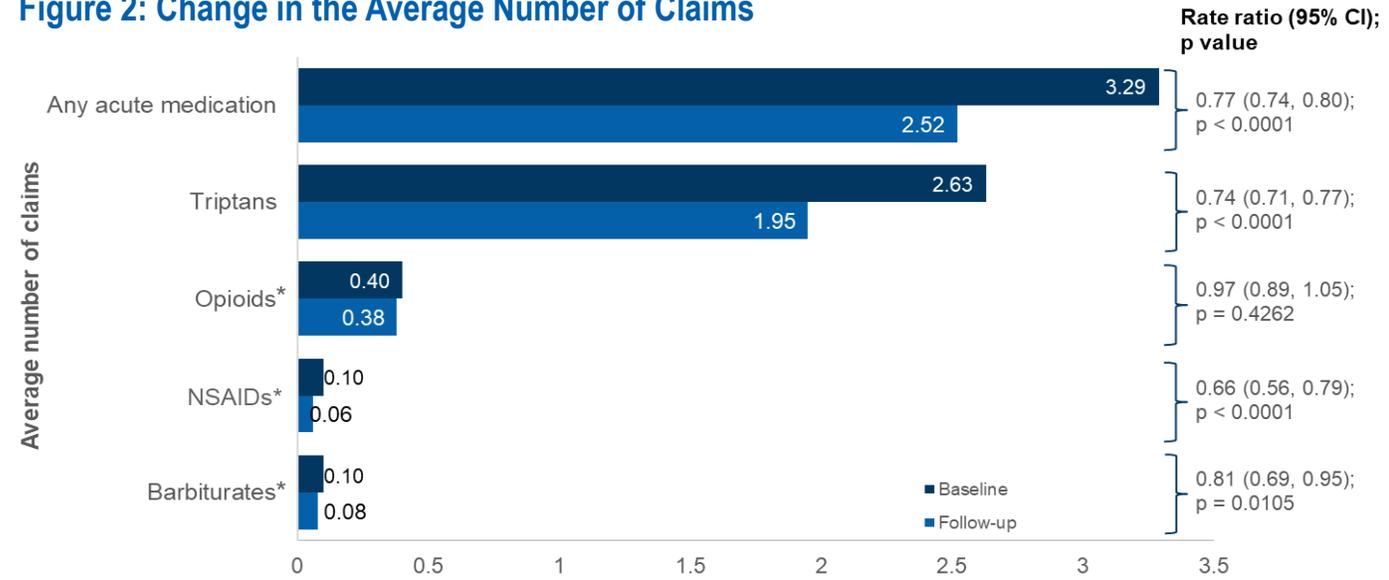
	Erenumab cohort (N = 3171)
Age at index date, mean (SD)	50.7 (13.6)
Female, n (%)	2689 (84.8)
CM w/o aura, n (%)	1982 (62.5)
<b>Index physician specialty, n (%)</b>	
Neurologist/headache specialist	2159 (68.1)
General practitioner	439 (13.8)
Nurse/physician assistant	293 (9.2)
Unknown/missing	154 (4.9)
<b>Region, n (%)</b>	
South	1536 (48.4)
West	684 (21.6)
Midwest	667 (21.0)
Northeast	284 (9.0)
<b>Insurance type, n (%)</b>	
Point of service (POS)	1569 (49.5)
Other	741 (23.4)
Health maintenance organization (HMO)	518 (16.3)
Exclusive provider organization (EPO)	214 (6.7)
Preferred provider organization (PPO)	129 (4.1)
<b>Selected comorbidities (&gt;10%), n (%)</b>	
Anxiety	1304 (41.1)
CV disease	1298 (40.9)
Depression	1295 (40.8)
Insomnia	731 (23.1)
Constipation	421 (13.3)

\*Use of non-migraine specific acute medications (NSAIDs, opioids, and barbiturates) required a migraine diagnosis on or before 7 days of the medication claim to proxy migraine-specific acute medication

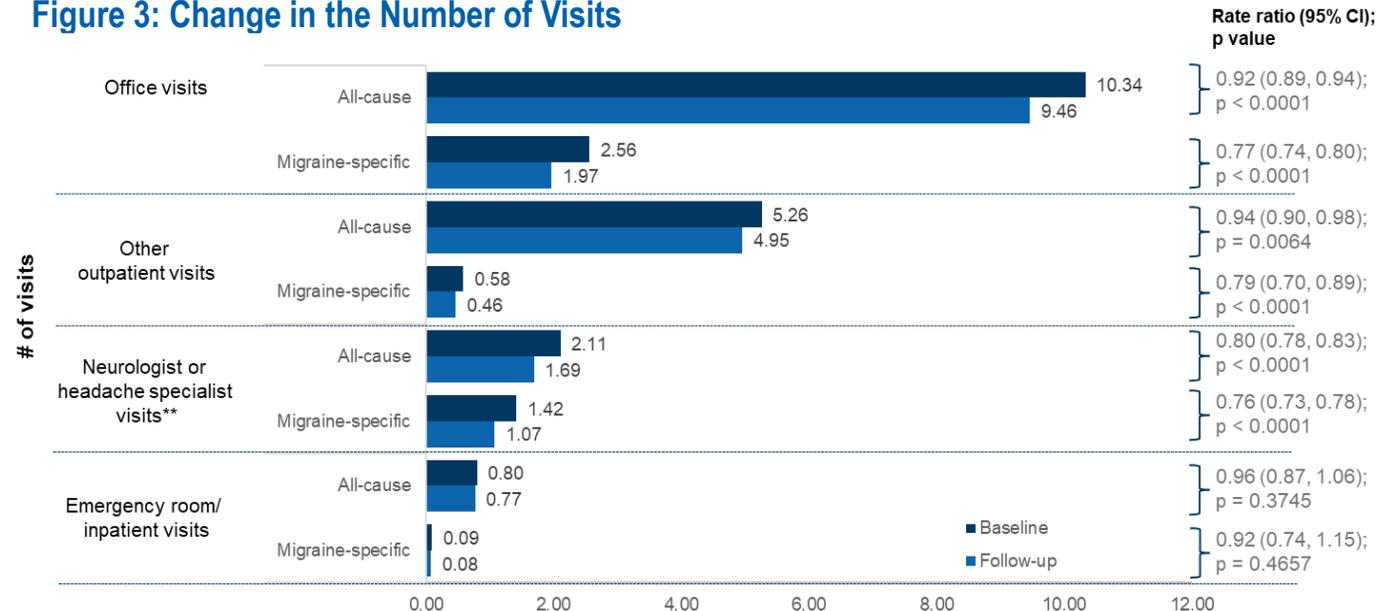
\*\*Neurologist or headache specialist visits are a subset of the office visits

CM, chronic migraine; CI, confidence interval; CV, cardiovascular; ER, emergency room; HCP, healthcare practitioner; n or #, number; NSAID, nonsteroidal anti-inflammatory drug; PS, propensity score; SD, standard deviation; SMD, standardized mean difference; w/o, without.

**Figure 2: Change in the Average Number of Claims**



**Figure 3: Change in the Number of Visits**

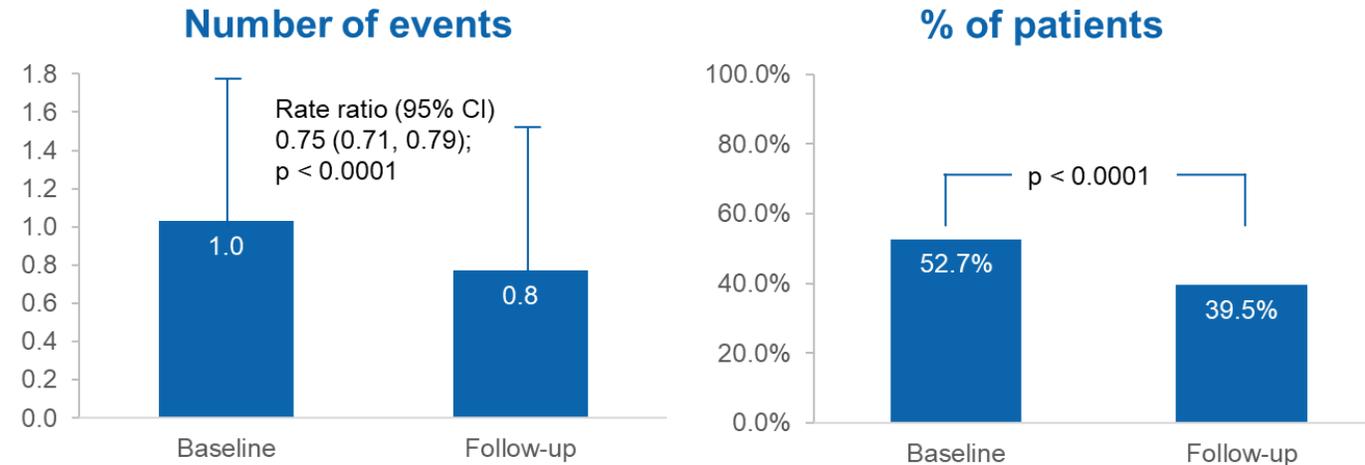




## Results and Conclusions

- For the composite outcome:
  - ⇒ Significant reduction in the mean (SD) number of events was observed from 1.03 (1.53) to 0.77 (1.48)
  - ⇒ Proportion of patients with any of the three events also significantly decreased (52.7% vs. 39.5%,  $P < 0.0001$ )

Figure 4: Composite Endpoint\*



\*Outpatient visit with a diagnosis of migraine and an associated acute medication claim, hospital admission or emergency room visit with a primary diagnosis for migraine. Any events occurred  $\leq 3$  days apart were counted only once. McNemar test for binary endpoint and Negative binomial model with repeated measure for count data were used.

## Conclusions

- Erenumab significantly reduces acute medication use and HCRU in real-world setting, hence significantly reducing the burden of the disease
- Significant reduction of 25% on the composite endpoint of the outpatient visits with an acute medication claim and migraine specific ER or IP visits shows the holistic benefit of erenumab in the real world
- A composite endpoint could be used as a proxy to evaluate migraine attacks, while further research is needed to validate the algorithm